

## TRACE ELEMENT ANALYSIS OF COAL BY NEUTRON ACTIVATION

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## INTRODUCTION

This report presents the technology methods used at the NASA Plum Brook reactor (PBR) to analyze coal by neutron activation analysis. The work was performed for the Environmental Protection Agency (EPA) Division of Air Surveillance at Research Triangle Park, North Carolina, by the NASA Plum Brook reactor at Sandusky, Ohio, under an interagency agreement. Unfortunately, the work was terminated in January 1973 when the Plum Brook reactor was closed.

The general scheme of analysis - sample preparation, irradiation, and sample counting - is described. The discussion of data reduction includes the computerized method, interference corrections, and the precision and accuracy of the method. Some typical trace element results are given for coal, fly ash, and bottom ash. The manpower requirement for the analysis of 1000 samples per year is stated.

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## THE ANALYSIS SCHEME

Figure 1 shows the overall analysis scheme used at PBRF to analyze coal and fly ash samples by neutron activation analysis (NAA). (These same techniques and methods, with only slight modification, were used for the NAA of kerosene, jet fuel, gasoline, fuel oil, residual oil, ore, air particulates on filters, bottom ash, sand, clam tissue, corn, cement, limestone, stack scrubber water, crab shells, and river water.)

The basic procedure used two aliquots of the sample. One aliquot was encapsulated in a polyethylene vial, the other in a synthetic quartz (Suprasil) vial. The polyethylene vial, containing 50 to 100 milligrams of coal (10 mg or less of fly ash), was irradiated for 5 minutes in a thermal neutron flux of  $1.5 \times 10^{14} n/cm^2/sec$ . Then the irradiated sample was counted at decay times of ~5 minutes, ~30 minutes, and 24 hours. The quartz vial was irradiated for 12 hours in the same flux and counted at about 3 weeks decay.

Samples were counted on a 4096 channel gamma ray spectrometer using a Ge (Li) detector with a crystal diameter of 35 mm and length of 27 mm. Acceptable counting distances ranged from 3 to 40 cm, with detector dead time restricted to <20% whenever possible. The detector resolution was 3 keV with linearity adjusted to 1 keV/channel and maintained at  $\pm 1$  channel or less.

## DATA REDUCTION

Altogether there were two irradiations associated with each pollution sample, and four counts - 5 minutes, 30 minutes, 24 hours, and 3 weeks decay. Each count of each aliquot produced a paper tape. Each tape, along with information regarding decay time, count time, flux level, sample weight, counting distance, and other parameters was processed through the data reduction code "SPECTRA."<sup>(1)</sup> Computing time on an IBM 360 Mod 67 was under 1 minute for all four tapes associated with one sample. Data were reported to EPA as parts per million of each trace element in the sample.

#### INTERFERENCE CORRECTIONS

EPA designates the elements Hg, Se, As, as "very hazardous." As a result, we devoted much attention to the accurate determination of these four elements by NAA. We found that, in coal, both selenium and ytterbium interfered with the detection of mercury; ytterbium interfered with selenium; and bromine and antimony interfered with arsenic.

The appendix to this paper describes how factorial experiment design techniques were used to derive empirical correction factors needed for the accurate determination of Hg, Se, and As.

#### PRECISION AND ACCURACY

Table 1 shows the standard deviation and the range of counting precision associated with the determination of trace elements in coal. The standard deviation is based on five aliquots.

Table 2 provides information on the accuracy of the NAA methods. Results of NAA of NBS standard reference materials (trace elements in a glass matrix) are compared with certified and interim NBS values. Other elements reported to EPA were frequently checked with homemade standards and with standards submitted by EPA.

#### TYPICAL TRACE ELEMENT RESULTS

Tables 3.1 to 3.4 show the form of computer outputs typically obtained, one for each of the four spectra associated with one sample. The data include the PBR sample number, the EPA identification number, and the input data required for the computerized data reduction. The third column shows the 56 elements that were routinely reported. Column four gives the results in ppm, and column 5 gives the standard deviation at 1 $\sigma$  in ppm.

Typical results are tabulated in table 4 for seven coal samples, bottom ash, and fly ash. The results are given in ppm. The elements are listed along with the isotopes actually detected. Examination of the data shows that calcium, cerium, iron, aluminum, barium, potassium, manganese, sodium, rubidium, tin, titanium, thorium, uranium, vanadium, and zirconium are concentrated in fly ash. An additional comment regarding uranium and thorium: these elements are naturally radioactive and are  $\alpha$ -emitters. For each ton of coal burned, the potential hazard exists of emitting 0.3 curies of  $\alpha$ -activity, based on 1 ppm of uranium in coal.

No element in table 4 shows a higher concentration in the bottom ash than in the fly ash.

#### THE ANALYSIS CAPABILITY

The analysis capability developed at the Plum Brook reactor was geared to analyze 1000 samples per year as a part time effort. The program was also geared to developing the technology to handle and irradiate a large variety of pollution related samples with a minimum of manpower. During a typical work week, 24 samples were irradiated, counted, and reported. Total manpower expended averaged 3.0 to 3.5 hours per sample. Computer running time amounted to approximately 1 minute per sample with 56 elements reported.

#### CONCLUDING REMARKS

The trace element analysis of coal using NAA has been shown to be an accurate, reliable, and instrumental method of analysis. The associated technology was also developed to permit the analysis of up to 56 trace elements in each of 1000 samples

per year as a part-time effort. Computerized data reduction reduced the total man-power required to 3.0 to 3.5 hours per sample.

#### APPENDIX - INTERFERENCE CORRECTIONS FOR MERCURY, SELENIUM, AND ARSENIC

by Anne Bodnar

The elements Hg, Se, As, and Cd are designated as "very hazardous" by EPA. This appendix describes the method used to derive the complex correction factors required to achieve an accurate determination of Hg, Se, and As by NAA. Cadmium did not require any special interference corrections.

We detected an accuracy problem with the determination of Hg in coal. A count at 5 to 6 weeks decay time produced a Hg result ranging from 2 to 10 times smaller than the value obtained at 3 weeks decay. A correction for Se interference on Hg was being made, but, because of the similarity in the half-lives of Hg<sup>203</sup> and Se<sup>75</sup>, the low Hg results at 6 weeks decay could not be explained.

A search of the Nuclear Data Tables<sup>(2)</sup> produced another interference: 4.2-day Yb<sup>175</sup>. Not only did the 282-keV peak of Yb<sup>175</sup> interfere with Hg<sup>203</sup>, but also the 400.7-keV peak of Se<sup>75</sup> interfered with the 396-keV peak area of Yb<sup>175</sup>, which was used for the Yb correction on Hg<sup>203</sup>. In addition, another Yb isotope, 32-day Yb<sup>169</sup> interfered with the 264-keV peak area of Se<sup>75</sup>, which is used in the correction on Hg<sup>203</sup> (peak area at 279 keV). These discrepancies were not eliminated by using theoretical corrections.

Finally, the problem was resolved by irradiating standards and mixtures of standards in a factorial experiment. The experiment design was a full factorial experiment with three variables (Hg, Se, Yb) at two levels, with replication, and with a center point added to test higher order effects. The high level was selected as 100 micrograms ( $\mu\text{g}$ ), the low level as 10  $\mu\text{g}$ . Table A-1 shows the treatments that were used.

Regression analysis on the data was used to estimate the coefficients in a predictive model equation. The dependent variable was chosen as the difference between the computer calculated value for Hg (or Se or Yb) and the true value. Independent variables were the other two elements plus plausible interactions (e.g., the interaction of Se-Yb on Hg). The coefficients derived for the predictive equations served as the basis for the empirical correction of Yb on Se and vice versa, and Yb-Se on Hg. Table A-2 compares the theoretical correction factors with the final form of the corrections based on the empirical data. A special subroutine was added to the SPECTRA computer program. The program was then tested by irradiating and analyzing other known samples. The calculated values agreed with the known sample contents. The method now allows us to determine Se and Hg in the presence of interferences which may be ten times greater in quantity.

This same technique has been used to determine arsenic in the presence of bromine and antimony.

#### REFERENCES

1. G. A. Borchardt, G. E. Hoagland, and R. A. Schmitt, *J. Radioanalyt. Chem.*, 6, 241 (1970).
2. Anon., "Nuclear Data Tables," Academic Press, New York, N.Y.

TABLE A-1

Treatment	Se	Hg	Yb	
1	-1	-1	-1	(-1) indicates low level, 10 $\mu\text{g}$
2	+1	-1	-1	
3	-1	+1	-1	(+1) indicates high level, 100 $\mu\text{g}$
4	-1	-1	+1	
5	+1	+1	-1	CP indicates $\frac{(\text{High level} + \text{low level})}{2}$
6	-1	+1	+1	
7	+1	+1	+1	
8	+1	-1	+1	
Replicates				
4	-1	-1	+1	
7	+1	+1	+1	
8	+1	-1	+1	
Center point	CP	CP	CP	

TABLE A-2

## 1. Theoretical corrections for Se and Yb interferences on Hg

$$1. A_c = A_u - 0.0369 X$$

$$2. B_c = B_u - 0.118 A_c$$

$$3. D_c = D_u - 0.959 B_c - 0.387 A_c$$

## 2. Empirical form of Se and Yb correction factors for interference on Hg

$$1. A_c = A_u - 0.0433 X$$

$$2. B_c = 0.443(B_u - 0.118 A_c)$$

$$3. D_c = D_u - 1.65 B_c - 0.387 A_c$$

where  $A_c$  = selenium 264.6 KeV area corrected for  $\text{Yb}^{169}$

$A_u$  = net area of selenium (264.6 KeV), uncorrected

$X$  = net area of  $\text{Yb}^{169}$  at 177.2 KeV

$B_c$  = ytterbium 396.1 KeV area corrected for selenium (264.6 KeV)

$B_u$  = net area of ytterbium (396.1 KeV) uncorrected

$D_c$  = mercury 279.1 KeV area corrected for selenium and ytterbium

$D_u$  = net area of mercury (279.1 KeV) uncorrected

TABLE 1. - PRECISION OF EPA ROUND ROBIN COAL SAMPLE

Element	ppm	% std dev ( $1\sigma$ )	Range of counting precision at 1 std dev, %
Ti	1 312	12	10-20
V	36	11	5-10
Al	15 700	9	0.6-1
S	<30 500	50	-----
U	0.98	8	8-12
Ba	340	12	5-8
Sr	93	10	8-11
I	2.8	14	12-30
Mn	38	7	0.5
Mg	890	28	12-33
Na	370	9	2-3
Cl	750	10	2
Ca	4 070	14	8-15
Cu	14	6	3-5
As	5.9	9	10-12
Br	20.	15	9-12
K	3 500.	10	3-4
Cd	<55	23	-----
Ce	17.3	5	1-2
Se	3.8	13	25-33
Hg	0.95	10	7-40
Cr	19	4	3-5
Cs	2.6	3	8-10
Ag	<0.4	25	-----
Zr	<70	55	-----
Zn	7.5	--	33
Fe	7 520	2	1
Co	5.5	15	1-13
Sb	6.4	24	8-15
Ni	<65	31	-----

TABLE 2. - COMPARISON OF NBS STANDARD REFERENCE MATERIALS WITH PBR RESULTS

	NBS 610*/PBR	NBS 612/PBR	NBS 614/PBR	NBS 616/PBR
Antimony	---	---	(1.06)/1.1±.1	0.078±.007/0.12±.02
Cerium	---	(39)/37±2	--	--
Cobalt	(390)/135±14	(35)/31±1	0.73±0.02/0.59±.006	--
Europium	---	(36)/26±1	0.99±.04/1.1±.6	--
Gold	(25)/20±2	(5)/4.7±1	(0.5)/1.0±.8	--
Lanthanum	---	(36)/35±15	0.63±.02/2	--
Thorium	---	37.6±.09/31.2±1	0.746±.007/0.58±.15	0.025±.004/0.018±.002
Scandium	---	--	0.59±.04/0.68±.23	0.026±.012/0.020±.004
Silver	(254)/180±80	22.0±.8/31±7	0.46±.02/0.57±.07	--

\* NBS values in parentheses are interim values for various reasons. Others are certified values.

TABLE 3.1 - TYPICAL INPUT AND OUTPUT DATA

SAMPLE NUMBER 2495  
---FPAFA1290

DECAY TIME	810.	SEC.
COUNT TIME	200.	SEC.
SAMPLE MASS	0.4400E-01	GRAMS
READ TIME	32.00	*
LENGTH OF IRRADIATION TIME	300.0	SEC.
FLUX LEVEL	0.1720E 15	NT. DPF SQ.CM. PDP CFR.
DISTANCE	10.0CM.	
AREA OF FILTER IRRADIATED	0.030	SQ.CM.
TOTAL AREA OF FILTER	0.000	SQ.CM.
VOLUME OF AIR FILTERED	0.00000000	FURTC METROS
KSPDCA	4	
RFC 2685 , ---FPAFA1290 ,	84	
PRC 2695 , ---FPAFA1290 ,	71	
RRC 2685 , ---FPAFA1290 ,	Y	
RRC 2695 , ---FPAFA1290 ,	A	
RFC 2495 , ---FPAFA1290 ,	S	

TABLE 3.2 - TYPICAL INPUT AND OUTPUT DATA

SCHOOL OF MINING 2637

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DECAY TIME	2775.	SEC.	
COUNT TIME	1000.	SEC.	
SAMPLE MASS	0.4400±.01	GRAMS	
DEAD TIME	20.00	?	
LENGTH OF IRRADIATION TIME	300.0	SEC.	
FLUX LEVEL	0.1720E 15	N.T.	DEP SEC.
DISTANCE	10.0CM.		
AFTER FILTER IRRADIATED	0.0000000	SEC.	RUBIC METERS
TOTAL AREA OF FILTER	0.0000000	SEC.	
VOLUME OF AIR FILTERED	0.0000000	SEC.	
KSCIDRA	4		
BRIG 2687	---EDAFAL1280	*	0.5753927E 00 *
BRIG 2687	---EDAFAL1280	*	0.5777538E 00 *
BRIG 2687	---EDAFAL1280	*	0.5299620E 02 *
BRIG 2687	---EDAFAL1280	*	0.3951313E 02 *
BRIG 2687	---EDAFAL1280	*	0.2574018E 01 *
BRIG 2687	---EDAFAL1280	*	0.24354018E 01 *
BRIG 2687	---EDAFAL1280	*	0.1126865E 04 *
BRIG 2687	---EDAFAL1280	*	0.1251408E 02 *
BRIG 2687	---EDAFAL1280	*	0.4522763E 01
BRIG 2687	---EDAFAL1280	*	0.4237236E 02 *
BRIG 2687	---EDAFAL1280	*	0.1403172E -01 *
BRIG 2687	---EDAFAL1280	*	0.22164131E 04 *
BRIG 2687	---EDAFAL1280	*	0.1333701E 04 *
BRIG 2687	---EDAFAL1280	*	0.76533624E -01
BRIG 2687	---EDAFAL1280	*	0.11211034E -01
BRIG 2687	---EDAFAL1280	*	0.6090129E 01
BRIG 2687	---EDAFAL1280	*	0.71572109E 01
BRIG 2687	---EDAFAL1280	*	0.5090911E 00
BRIG 2687	---EDAFAL1280	*	0.6813162E -01
BRIG 2687	---EDAFAL1280	*	0.6419050E 01
BRIG 2687	---EDAFAL1280	*	0.2024641E 02

TABLE 3.3 - TYPICAL INPUT AND OUTPUT DATA

SAMPLE NUMBER 2654  
---EPAFA12R0

	DECAY TIME	4800. SEC.	
	COUNT TIME	1000. SEC.	
	SAMPLE MASS	0.4400E-01 GRAMS	
	DEAD TIME	10.00 SEC.	
	LENGTH OF IRRADIATION TIME	300.0 SEC.	
	FLUX LEVEL	0.1720E-15 NT. DEG SQ.CM. PER SEC.	
	DISTANCE	3.0CM.	
	AREA OF FILTERED IRRADIATION	0.000 SQ.CM.	
	TOTAL AREA OF FILTERED	0.000 SQ.CM.	
	VOLUME OF AIR FILTERED	0.00000000 CURIC METERS	
K SUPRA			
PFC 2604	----EPAFA12R0	SM	0.49226C21F 00 *
PNC 2654	----EPAFA12R0	F1	0.16403475F 00 *
PFC 2604	----EPAFA12R0	F2	0.12648592E 01 *
PFC 2594	----EPAFA12R0	A11	0.21256555E 00 *
PNC 2654	----EPAFA12R0	F11	0.72881180E 01 *
PFC 2694	----EPAFA12R0	A5	0.1A097029E 01 *
PFC 2694	----EPAFA12R0	ap	0.346650284F 02 *
PFC 2694	----EPAFA12R0	4	0.49226C21F 00 *
PNC 2694	----EPAFA12R0	GA	0.1745166RF 01 *
PFC 2694	----EPAFA12R0	K	0.21976392E 03 *
PNC 2654	----EPAFA12R0	L4	0.4907530RF 01 *
PFC 2604	----EPAFA12R0	PT	E < 0.90368561F 02 *
PFC 2694	----EPAFA12R0	W	E < 0.14073496E 01 *
PNC 2604	----EPAFA12R0	GD	E < 0.1A51574E 02 *
PNC 2694	----EPAFA12R0	CD	E < 0.50213211E 02 *
PNC 2654	----EPAFA12R0	QE	E < 0.97890139E-01 *
PNC 2604	----EPAFA12R0	MN	E < 0.65644312F 03 *

TABLE 3.4 - TYPICAL INPUT AND OUTPUT DATA

SAMPLE NiIJwafq  
27C1  
---EPFA1280

	DECAY TIME	2160000.	SEC.
COUNT TIME	4000.	SEC.	
SAMPLE MASS	0.4085F 00	GRAMS	
DEAD TIME	13.00	%	
LENGTH OF IRRADIATION TIME	43270.0	SEC.	
FLUX LEVEL	0.110E 16	NT. DEG SQ.CM. PER SEC.	
DISTANCE	3.0CM.		
AREA OF FILTER IRRADIATED	0.000	SQ.CM.	
TOTAL AREA OF FILTERED	0.000	SQ.CM.	
VOLUME OF AIR FILTERED	0.00000000	CUBIC METERS	
KILODRA 1			
PRG 2701	----EPFA1280	CC	
PRG 2701	----EPFA1280	L1	
PRG 2701	----EPFA1280	SC	
PRG 2701	----EPFA1280	HG	
PRG 2701	----EPFA1280	TR	
PRG 2701	----EPFA1280	TH	
PRG 2701	----EPFA1280	CR	
PRG 2701	----EPFA1280	HE	
PRG 2701	----EPFA1280	ND	
PRG 2701	----EPFA1280	ZR	
PRG 2701	----EPFA1280	CS	
PRG 2701	----EPFA1280	SC	
PRG 2701	----EPFA1280	RR	
PRG 2701	----EPFA1280	7N	
PRG 2701	----EPFA1280	TA	
PRG 2701	----EPFA1280	FE	
PRG 2701	----EPFA1280	CO	
PRG 2701	----EPFA1280	SR	
PRG 2701	----EPFA1280	AC	
PRG 2701	----EPFA1280	SN	
PRG 2701	----EPFA1280	NI	
PRG 2701	----EPFA1280	YR	
	0.72096167F 01	*	0.10686743E 00
	0.17842479F 00	*	0.13465714E-01
	0.25705E0CE 01	*	0.64358050E 00
	< 0.64539255F 00	*	0.12052262E 00
	0.12754770E-01	*	0.62859185E-03
	0.95413691F 00	*	0.14910755F-01
	0.93224478E 01	*	0.27451810F 00
	0.30085230F 00	*	0.18846393E-01
	< 0.27277546F 01	*	0.13943005F 01
	0.35244324F 02	*	0.14130380E 02
	0.40247416E 00	*	0.54329596E-01
	0.16070574F 01	*	0.52692741E-02
	0.32748070F 01	*	0.60219342F 00
	-0.35999384F 01	*	-0.10403924E-01
	0.10774275E 00	*	0.20200126F-01
	0.26460225F 04	*	0.22373123F 02
	0.35183659F 01	*	0.29760309E-01
	0.9539272AE 00	*	0.92135251E-01
	< 0.13639234F 00	*	0.96802831E-01
	0.16124039E 02	*	0.42969913F 01
	0.33774506F 02	*	0.12184061E 02
	0.76171124F 00	*	0.13029099E 00

TABLE 4. - ENVIRONMENTAL PROTECTION AGENCY SAMPLES\* - COAL, FLY ASH, AND BOTTOM ASH

Element / isotope	Coal L-14	Coal L-15	Coal L-16	Coal L-17	Coal L-18	Coal L-19	Coal L-20	Bottom ash L-21	Fly ash L-22	Fly ash L-23
Ag110M	<0.9	<0.7	<0.6	<0.6	<0.2	<0.2	<0.3	<0.2	<0.4	<2
Al28	17 200	8 080	13 300	12 100	10 800	6 170	27 600	67 500	89 600	99 700
As76	7.00	<1.2	4.8	<5	2.44	1.6	<6	<5	<7	<6
Au198	0.003	C.10	0.07	0.15	0.06	0.02	<0.4	<0.5	<2	<2
Be139	92	34	48	53	45	29	565	1500	1641	2370
Br82	20	3	22	19	9.0	16	<20	<15	<50	<30
Ca49	7 740	3 640	5640	<5 000	3290	1910	5960	14 300	23 200	26 200
Cd115	<300	<110	<110	<90	<200	<30	<100	<900	<1200	<2000
Ce141	21.1	8.83	16.2	13.0	10.6	8.97	28.0	71.8	104	103
Cf138	2820	1220	2160	1450	1250	2350	189	94	<200	168
Ni(Co58)	<90	<30	<80	<70	<60	<30	<20	<50	<30	<50
Co60	5.98	3.07	5.22	4.40	3.84	4.02	1.92	4.72	6..6	6.8
Cr51	24.2	18.0	21.4	19.4	19.3	12.2	5.26	14.8	22	21
Cs134	3.32	1.56	2.58	2.35	2.03	1.13	0.92	2.81	4.4	4.1
Cu64	<30	29	<20	<20	<20	<20	<50	37	57	70
Dy165	1.2	0.58	0.76	0.77	0.67	0.42	0.90	2.0	2.8	3.50
Eu152m1	0.56	0.2	0.32	0.31	.30	.18	.36	0.91	1.2	1.5
Fe59	8970	13 700	9500	10 900	11 600	4550	6150	20 000	24 400	23 600
Ga72	7.2	<2	4.0	4.2	5.5	2.9	<6	<2	<30	<30
Ge75	<70	<4	<120	<40	<150	<150	<20	<200	<300	<400
Hf181	1.05	0.50	0.81	0.65	0.60	0.42	2.28	6.50	8.5	8.7
Hg203	<0.7	<0.5	<0.6	1.91	0.16	<0.3	<0.5	<0.7	<0.9	<0.9
Gd159	<70	<60	<40	<30	<40	<20	<80	<120	<140	<400
I128	<0.2	<1	<0.5	1.8	<2	0.65	<2	<1	<2	<20
In116m1	<0.05	<0.03	<0.02	0.073	0.029	<0.01	<0.05	<0.07	<0.02	<0.07

\*The EPA could not reveal the source of these samples.

TABLE 4. - Continued. ENVIRONMENTAL PROTECTIVE AGENCY SAMPLES - COAL, FLY ASH, AND BOTTOM ASH

Element/ isotope	Coal L-14	Coal L-15	Coal L-16	Coal L-17	Coal L-18	Coal L-19	Coal L-20	Bottom ash L-21	Fly ash L-22	Fly ash L-23
Ir <sup>192</sup>	5.3	1.9	2.2	6.7	1.6	1.8	7	11	9.8	1.3
K <sup>42</sup>	7130	2464	3932	3748	3420	1690	1810	7170	9064	9800
La <sup>140</sup>	17.2	5.7	7.2	26	7.3	5.4	20	29	25	57
Lu <sup>177</sup>	0.58	0.36	0.42	0.32	0.35	0.22	0.51	1.23	1.6	1.8
Mg <sup>27</sup>	<450	<850	<1200	<4000	585	<600	1320	<11 000	2400	<9000
Mn <sup>56</sup>	95	53	62	49	38	25	93	300	371	422
Mo <sup>99</sup>	<600	<700	<260	<350	<300	<200	<3100	<2200	<13 000	<6000
Na <sup>24</sup>	1250	882	1070	833	757	487	2850	6080	8430	14 000
Nd <sup>147</sup>	<7	<2	<2	<5	<7	<3	<7	<10	<40	<30
Pt <sup>197</sup>	<300	<170	<160	<230	<200	<100	<1000	<700	<1400	<3200
Rb <sup>86</sup>	25.6	12.3	20.7	21.4	16.8	7.98	8.66	15.9	45	41
Re <sup>188</sup>	<0.3	<0.2	<0.2	<0.3	<0.4	<0.3	<0.5	<2	<1	<0.4
Rh <sup>104</sup>	<0.03	<0.09	<0.04	<0.02	<0.06	<0.06	<0.07	<0.3	<4	<0.3
S <sup>37</sup>	*	37 500	<53 000	<7 000	<9000	<30 000	<200 000	168 000	<200 000	<300 000
Sc <sup>46</sup>	4.56	2.67	3.84	3.42	3.17	2.07	3.02	7.42	11.4	11.7
Sb <sup>124</sup>	1.39	0.84	1.12	0.93	0.62	0.86	0.80	0.50	2.1	1.8
Se <sup>75</sup>	4.15	3.12	2.99	3.19	2.86	1.77	3.07	2.10	5.98	5.72
Sm <sup>153</sup>	2.23	0.82	1.2	1.2	1.1	0.81	1.5	3.5	4.4	4.9
Sr <sup>87m</sup>	56	<30	36	<20	<40	<20	106	<100	223	<200
Sn <sup>117m</sup>	76.5	<40	74.2	66	50	34	86	428	555	422
Ta <sup>182</sup>	-	-	-	-	-	-	-	-	-	-
Tb <sup>160</sup>	0.35	0.12	0.27	0.25	0.18	0.17	0.65	1.66	2.4	2.4
Th(Pa <sup>133</sup> )	0.036	0.018	0.027	0.025	0.022	0.015	0.048	0.12	0.18	0.18
	4.19	2.16	3.23	2.84	2.84	1.73	7.49	19.1	25	25

\* Data missing.

TABLE 4. - Concluded. ENVIRONMENTAL PROTECTIVE AGENCY SAMPLES - COAL, FLY ASH, AND BOTTOM ASH

Element/ isotope	Coal L-14	Coal L-15	Coal L-16	Coal L-17	Coal L-18	Coal L-19	Coal L-20	Bottom ash L-21	Fly ash L-22	Fly ash L-23
Tl <sup>205</sup>	1680	912	608	980	908	529	893	3820	4280	3428
U <sup>239</sup>	1.5	0.96	0.82	1.14	0.82	0.68	1.3	3.7	4.7	4.9
V <sup>52</sup>	34	23	26	27	23	15	18	29	65	64
W <sup>187</sup>	<4	<0.3	<0.4	<2	<2	<2	<20	<15	<20	<30
Yb <sup>175</sup>	3.68	1.88	4.06	1.83	0.46	0.91	0.84	2.36	3.8	2.9
Zn <sup>65</sup>	264	<3000	220	148	118	<400	<2000	187	299	361
Zr <sup>95</sup>	<50	<50	<50	<40	<50	<30	107	261	400	556

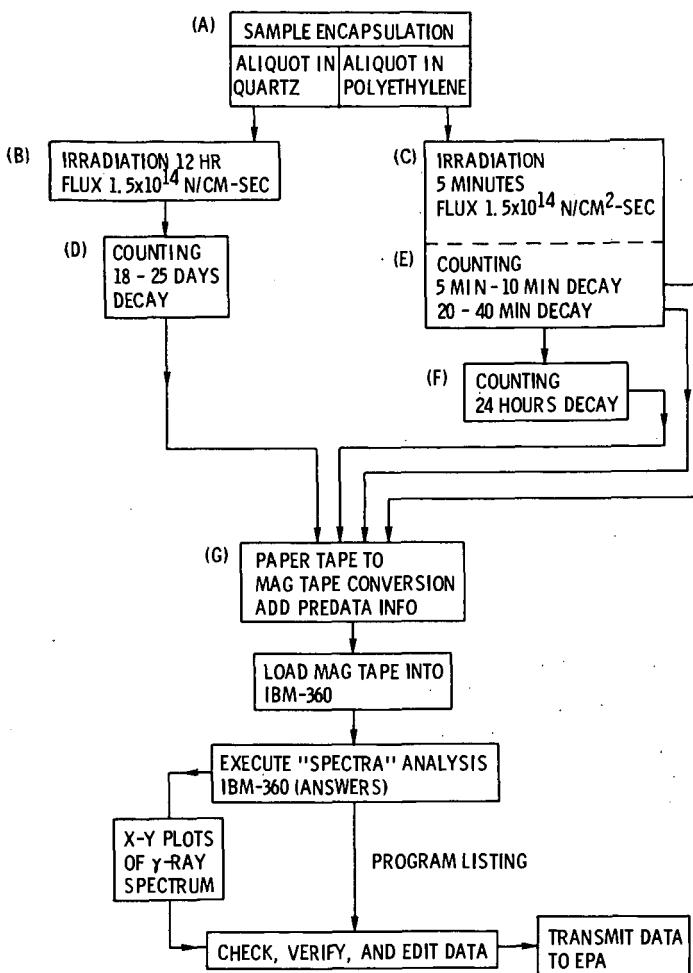


Figure 1. - Flow chart showing scheme for sample irradiation, counting, and data reduction of coal samples.